Under the Paperwork Reduction Act of 1995, no persons are required to

INFORMATION DISCLOSURE
STATEMENT BY APPLICANT
(Not for submission under 37 CFR 1.99)

Application Number		10541987			
Filing Date		2005-07-11			
First Named Inventor Dolf		H. J. Van Casteren, et al			
Art Unit		2821			
Examiner Name					
Attorney Docket Number		NL030036	_		

				Attorney	y Doc	ket Number		NL030036			
					U.S.I	PATENTS				Remove	
Examiner Initial*	Cite No	Patent Number	Kind Code ¹	Issue Dat	to	Name of Patentee or Applicant of cited Document		Pages,Columns,Lines when Relevant Passages or Rele Figures Appear		re van	
	1	6144171	A	2000-11-0	07	DEURLO	ÞΕ	TAL			
	2	4952819	A	1990-08-2	28	HERRMAI	NN	DIETER			
If you wish	to a	dd additional U.S. Pate	nt citatio	n informati	ion pl	ease click the	Ac	id button.		Add	
			U.S.P	ATENT A	PPLIC	CATION PUB	LIC	CATIONS		Remove	
Examiner Initial*	Cite No	Publication Number	Name of Patentee or Application Code! Date Name of Code Document			Releva	s,Columns,Lines whe ant Passages or Rele es Appear	re van			
	1										
If you wish	n to a	dd additional U.S. Publ	ished Ap	plication c	itation	n information	ple	ase click the Add	d buttor	n. Add	
				FOREIGN	I PAT	ENT DOCUM	(E)	VTS		Remove	
Examiner Initial*	Cite No	Foreign Document Number ³	Country Code ²			Publication Date	A	lame of Patentee opplicant of cited occument	or	Pages, Columns, Line where Relevant Passages or Relevar Figures Appear	7.
	1										
lf you wish	n to a	dd additional Foreign P							button		_
			NON	I-PATENT	LITE	RATURE DO	CL	JMENTS		Remove	

	Application Number		10541987	
INFORMATION DIGGS COURT	Filing Date		2005-07-11	
INFORMATION DISCLOSURE STATEMENT BY APPLICANT	First Named Inventor	Dolf I	H. J. Van Casteren, et al	
(Not for submission under 37 CFR 1.99)	Art Unit		2821	
(······	Examiner Name			
	Attorney Docket Numb	er	NL030036	

Examine Initials*	Cite No	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc), date, pages(s), volume-issue number(s), publisher, city and/or country where publisher.	Ţ5	
	1			

If you wish to add additional non-patent literature document citation information please click the Add button Add

EXAMINER SIGNATURE

Examiner Signature | Date Considered | Date Considered |

"EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through a citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

1 See Kind Codes of USPT/D Patient Documents at year USPT(D,QQ/ or MEEP 801.04. ² Enter office that issued the document, by the bit-office code (WIPD Standard ST.3.) ² Sort disparses patient colorments, by excitation of the year of the Emperor most procedure the sent number of the patient colorment.

*Kind of document by the opportuite symbols as indicated on the document under WIPD Standard ST.16 if possible. ³ Applicant is to place a check mark here if English tanguage brantation is attached.

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		10541987
Filing Date		2005-07-11
First Named Inventor Dolf I		H. J. Van Casteren, et al
Art Unit		2821
Examiner Name		
Attorney Docket Number		NL030036

CERTIFICATION STATEMENT

Please see 3'	7 CFR 1.97	and 1.98 to make	the appropriate selection(s):
---------------	------------	------------------	-------------------------------

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. Sec 37 CFR 197(eV1).

ΩR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquity, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 156(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 197(4)(2)

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.
- 7 None

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

_						
Signature	/R.J.KRAUS/	Date (YYYY-MM-DD)	2006-09-11			
Name/Print	Robert J. Kraus	Registration Number	26358			

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is for life and by the USPTO to process) an application. Confidentiality is governed by \$5 U.S. C.12 and 3T CFR.

1.14. This collection is estimated to take I hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, u.S. Operatment of Commence, P. O. Box 1430, Alexandriu, V.S. 2213.1450, DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, V.A. 2213.1450.

Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that (1) the general authority for the collection of this information is \$3 U.S.C. 2(b)(2); (2) famishing of the information solicided is voluntary, and (3) the principal purpose for which the information is used by the U.S. Patient and Trademan Kollie is to process and/or examine your submission related to a patient application or patient. If you do not furnish the requested process and/or examine your submission related to a patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested to the patient process and/or examine your submission, which may related that the patient process and/or examine your submission, which may

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
 - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement neodiations.
 - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record partains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
 - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974. as amended, pursuant to 5 U.S.C. 552a(m.).
 - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
 to the Patent Cooperation Treaty.
 - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
 - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uturing an inspection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 44 U.S.C. 2504 and 2506. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make relevant for the state of the s
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S. C. 12(b) or issuance of a patient pursuant to 35 U.S. C. 15.1 Further, a record may be disclosed, subject to the limitations of 37 CFR.114, as a routine use, to the public if the record via flori of mapplication which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inseptions or an issued patient.
 - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.